

--2. (Amended) The process [Process] according to claim, wherein the expressed retroviral mRNA is specifically detected by nucleic acid amplification using primers, one of which is specific for the poly(A) signals present in the 3' R-poly(A) sequence at the 3' extremity of the retrovirus.--

--3. (Amended) The process [Process] according to claim 1, wherein the protein or peptide expressed by the endogenous retrovirus is detected using antibodies specific for the said retroviral protein or peptide.--

--4. (Amended) The process [Process] according to claim 1, wherein the antibodies specific to retroviral protein are detected by use of the retroviral protein, or fragments thereof with which the antibodies specifically react.--

--5. (Amended) The process [Process] according to claim 1, wherein SAg activity specifically associated with said HERV is detected, the biological sample being a biological fluid containing MHC Class II cells or cells induced to express MHC Class II molecules, this sample being contacted with cells bearing one or more variable (V)- β T-cell receptor chains, and detecting preferential proliferation of the V β subset, or one of the v β subsets characteristic of said autoimmune disease.--

--6. (Amended) The process [Process] according to claim 1, wherein the autoimmune disease is type I diabetes and the associated retrovirus having SAg activity is IDDMK_{1,2} 22 comprising the 5' long terminal repeat shown in Figure 7A, the 3' short terminal repeat shown in Figure 7B, or the env encoding sequence shown in Figure 7C, Figure 7D or Figure 7E, or variants thereof presenting approximately at least 90% sequence identity.--

--7. (Amended) The process [Process] according to claim 6, wherein the expressed retroviral RNA is specifically detected by nucleic acid amplification using primers, one of which is specific for poly(A) signals present in the 3' R-poly (A) sequences at the 3' extremity of IDDMK_{1,2}22.--

~~-8. (Amended) The process [Process] according to claim 7, wherein the poly (A) specific primer is~~

~~5' TTTTGTAGTCCCCTTAGTATTTATT 3' (SEQ ID NO: 26) or
5' T₍₂₀₎GAGTCCCCTTAGTATTTATT 3' (SEQ ID NO: 49)--~~

~~--9. (Amended) The process [Process] according to claim 6, wherein protein expressed by IDDMK_{1,22} is detected, said protein being either the protein encoded by the N-terminal moiety of the env coding region of IDDMK_{1,22} as illustrated in Figure 7D or 7G, or the protein encoded by the pol coding region, as illustrated in Figure 7H, or a protein having at least 90% homology with the illustrated protein, or a fragment of said proteins having at least 6 amino-acids.--~~

~~--10. (Amended) The process [Process] according to claim 6, wherein antibodies specific for env or pol proteins expressed by IDDMK_{1,22} are detected using the env or pol proteins illustrated in Figure 7D, 7G or 7H, or a protein having at least 90% homology with the illustrated protein, or a fragment of said proteins having at least 6 amino-acids--~~

~~--11. (Amended) A Human endogenous retrovirus having superantigen activity, and being associated with human autoimmune disease, said retrovirus being obtainable from RNA prepared from a biological sample originating from a human autoimmune source, by carrying out the following steps:~~

- ~~i) isolating [isolation of] the 5' R-U5 ends of expressed putative retroviral genomes using nucleic acid amplification, the 3' primer being complementary to known "primer binding sites" (pbs) and the 5' primer~~

004370-000000
Smit

- being an oligonucleotide anchor;
- ii) isolating [isolation of] the 3' R-poly(A) ends corresponding to the 5' R-U5 ends, by use of primers specific for the R regions isolated in step i);
 - iii) amplifying [amplification of] the conserved RT-RNase H region within the pol gene by using degenerate primers corresponding to the conserved region;
 - iv) amplifying [amplification of] the 5' moiety of the putative retroviral genome by using primers specific for the different U5 regions isolated in step i) in conjunction with a primer specific for the 3' end of the central pol region isolated in step iii);
 - v) amplifying [amplification of] the 3' moiety of the putative retroviral genome using primers specific for the central pol region isolated in step iii) in conjunction with primers specific for the poly(A) signals present in the 3' R-poly(A) sequences isolated in step ii); and
 - vi) confirming [confirmation of] the presence of an intact retroviral genome by amplification using primers specific for its predicted U5 and U3 regions.--

--12. (Amended) A proviral [Proviral] DNA of a retrovirus according to claim 11.--

--13. (Amended) A proviral [Proviral] DNA according to claim 12 obtainable from a biological sample of human origin by:

- i) obtaining retroviral RNA according to the method of claim 11, and further,
- ii) generating a series of DNA probes from the retroviral RNA obtained in i);
- iii) hybridising under stringent conditions, the probes on a genomic human DNA library;

iv) isolation of the genomic sequences hybridising with the probes.--

--14. (Amended) A nucleic [Nucleic] acid molecule comprising fragments of the retroviral RNA or DNA according to claim 11 [any one of claims 11 to 13], said fragment having a length of at least 15 nucleotides and preferably at least 30 nucleotides.--

--15. (Amended) A nucleic [Nucleic] acid molecule according to claim 14, encoding SAg activity of the retrovirus.--

--16. (Amended) A nucleic [Nucleic] acid molecule according to claim 15 derived from an endogenous human retrovirus open reading frame and optionally containing at least one internal stop codon.--

--17. (Amended) A nucleic [Nucleic] acid molecule according to claim 15 [or 16] comprising the retroviral env gene.--

--18. (Amended) A nucleic [Nucleic] acid molecule comprising a sequence complementary to the nucleic acid molecule of claim 11 [molecules of any one of claims 11 to 17].--

--19. (Amended) A nucleic [Nucleic] acid molecule according to claim 18 comprising a ribozyme or antisense molecule to a human retrovirus having SAg activity to a proviral DNA of said retrovirus or a fragment thereof.--

--20. (Amended) A nucleic [Nucleic] acid molecule capable of hybridizing in stringent conditions, with the nucleic acid molecules of claim 11 [any one of claims 11 to 19].--

004206460
AA
crit

--21. (Amended) A vector [Vector] comprising a nucleic acid molecules of claim 11 [any one of claims 11 to 20].--

--22. (Amended) A nucleic [Nucleic] acid molecule comprising at least one of the sequences illustrated in Figures 7A, 7B, 7C, 7D, 7E, or a nucleic acid sequence encoding the POL protein shown in Figure 7H, or a sequence exhibiting at least 90% homology with any of these sequences , or a fragment of any of these sequences having at least 20 nucleotides, and preferably at least 40 nucleotides.--

--23. (Amended) A nucleic [Nucleic] acid molecule having a sequence at least partially complementary to the sequence of any of the nucleic acid molecules [sequences] according to claim 22.--

--24. (Amended) A nucleic [Nucleic] acid molecule according to claim 22 comprising a ribozyme or antisense.--

--25. (Amended) A nucleic [Nucleic] acid molecule which is HERV IDDMK_{1,2-22} comprising each of the sequences illustrated in Figures 7A, 7B, 7C, or sequences having at least 90% identity with these sequences, having a size of approximately 8.5 kb, having SAg activity encoded within the **env** region illustrated in Figure 7D or 7E, said SAg activity being specific for V β 7- TCR chains.--

--26. (Amended) A protein [Protein] or peptide having at least 6 amino acids, characterised in that:

- it exhibits SAg activity and optionally is capable of giving rise, directly or indirectly, to autoreactive T-cells targeting tissue characteristic

Accepted
for publication
10/10/98

- it is encoded by a human endogenous retrovirus; and
- it is obtainable from biological samples of patients having autoimmune disease.--

--27. (Amended) A protein [Protein] or peptide according to claim 26, encoded by the env gene of the HERV, or a portion thereof.--

--28. (Amended) A protein [Protein] or peptide according to claim 27 corresponding to a protein or peptide resulting from a premature translational stop, and/or from a frame shift in the translation of a retroviral open reading frame.--

--29. (Amended) A protein [Protein] or peptide [according to any one of claims 25 to 28] obtainable by introducing viral DNA of claim 13 or fragments thereof, or corresponding synthetic DNA into a eukaryotic cell under conditions allowing the DNA to be expressed, and recovering said protein.--

--30. (Amended) A protein [Protein] according to claim 26 [any one of claims 26 to 29] comprising the amino acid sequence shown in Figure 7D, Figure 7F, Figure 7G, Figure 7H, or an amino acid sequence having at least 80% and preferably at least 90% homology with the illustrated sequences, or a fragment of said sequence having at least 6 amino acids.--

--31. (Amended) An antibody [Antibodies] capable of specifically recognising a protein or peptide according to claim 26 [any one of claims 26 to 30].--

--32. (Amended) An antibody [Antibodies] according to claim 31 which is [are] monoclonal.--

--33. (Amended) An antibody [Antibodies] according to claim 31 [or 32] which specifically recognises [recognise] a HERV protein having SAg activity and which has [have] the capacity to block SAg activity.--

--34. (Amended) A cell-line [Cell-line] transfected with and expressing a human retrovirus or a portion thereof or a nucleic acid molecule according to claim 11 [any one of claims 11 to 25].--

--35. (Amended) A non-human [Non-human] cells transformed with and expressing a human retrovirus or a nucleic acid molecule according to claim 11 [any one of claims 11 to 25].--

--36. (Amended) A cell-line [Cell-line] according to claim 34 [or 35] said cell lines or cells being MHC Class II⁺ and expressing a protein having SAg activity.--

--37. (Amended) A process [Process] for identifying substances capable of binding to retroviral protein or peptide according to claim 26 [any one of claims 26-30], comprising contacting the substance under test, optionally labelled with detectable marker, with the said retroviral protein or peptide having SAg activity, and detecting binding.--

--38. (Amended) A process [Process] for identifying substance capable of blocking SAg activity of an endogenous retrovirus associated with autoimmune disease, comprising introducing the substance under test into an assay system

00427663
A2
Cm. +

--39. (Amended) A process [Process] according to claim 38, wherein the cells bearing V β -T cell receptor chains are T-cell hybridoma and V β -specific stimulation is determined for example by measurement of IL-2 release, or measurement of T-cell proliferation.--

--40. (Amended) A process [Process] according to claim 38 [or 39,] comprising an additional preliminary screening step for selecting substances capable of binding to retroviral protein having SAg activity [,said screening step being according to claim 38].--

--41.(Amended) A process [Process] for identifying substances capable of blocking transcription or translation of human endogenous retroviral (HERV) SAg-encoding nucleic acid sequences, said SAg being associated with a human autoimmune disease, comprising:

- i) contacting the substance under test with cells expressing endogenous retroviral protein or peptide having SAg activity, according to one of the claims 26 to 30 and
- ii) detecting loss of SAg protein expression using SAg protein markers such as specific, labelled anti-SAg antibodies.--

--42. (Amended) A process [Process] according to claim 41, wherein the cells expressing HERV protein having SAg activity are MHC Class II⁺ cells, and the process further comprises detection of loss of SAg activity by the process of claim 38.--

--43. (Amended) A kit [Kit] for screening substances capable of blocking SAg activity of a retrovirus associated with an autoimmune disease, or of blocking transcription or translation of the retroviral SAg protein, comprising:

- MHC Class II⁺ cells transformed with and functionally expressing said retroviral SAg
- cells bearing V β T-cell receptor chains of the family or families specifically stimulated by the HERV SAg;
- means to detect specific V β stimulation by HERV SAg;
- optionally, labelled antibodies specifically binding to the retroviral SAg.--

--44. (Amended) A protein [Protein] or peptide derived from a retroviral SAg according to claim 26 wherein the [protein is modified so as to be devoid of SAg activity and is capable of generating a immune response against SAg, involving either antibodies and/or T-cell response.--

--45. (Amended) A protein [Protein] according to claim 44, whereing the modification consists of denaturation, or of a truncation, or of a deletion, insertion or replacement mutation of the SAg protein.--

--46. (Amended) A protein [Protein] according to claim 44 [or 45] for use as a prophylactic or therapeutic vaccine against autoimmune disease associated with retroviral SAg.--

--47. (Amended) A vaccine [Vaccine] comprising an immunogenically effective amount of a protein according to claim 44 [or 45] in association with a pharmaceutically acceptable carrier and optionally adjuvant.--

--48. (Amended) A nucleic [Nucleic] acid molecule encoding human retroviral SAg according to claim 15 or a modified form of said molecule for use as a prophylactic or therapeutic DNA vaccine against autoimmune disease associated with the retroviral SAg.--

--49. (Amended) A substance [Substance] identifiable by the process according to claim 37 [any one of claims 37 to 42] for use in therapy and/or prevention of autoimmune disease associated with the HERV SAg.--

--50. (Amended) A use [Use] of substance capable of inhibiting retroviral function for the preparation of a medicament for use in therapy and/or prevention of autoimmune disease associated with retroviral SAg.--

--51. (Amended) Use according to claim 50, wherein the substance capable of inhibiting retroviral function is Azido Deoxythymidine (A.Z.T.).--

--53. (Amended) A process [Process] for detecting human autoimmune disease associated with expression of human endogenous retrovirus Superantigen (SAg), said process comprising at least one of the following steps:

- i) detecting the presence of any expressed retrovirus in a biological sample of human origin; and
- ii) detecting the presence of SAg activity in a biological sample of human origin containing MHC

A3
cncld.

03473:004926

A3
Cm.t

--54. (Amended) A process [Process] according to claim 53, wherein the expressed retrovirus is detected by detection of reverse transcriptase activity.--

--55. (Amended) A process [Process] according to claim 54, wherein the expressed retrovirus is detected by carrying out nucleic acid amplification reaction on RNA prepared from the biological sample, using as 3' primer a sequence complementary to known retroviral «primer binding sites» (pbs), and as 5' primer a non-specific anchor sequence.--

--56. (Amended) A process [Process] according to claim 53, wherein the presence of SAg activity is detected by contacting the biological sample containing MHC Class II⁺ cells with cells bearing one or more variable (V)- β T-cell receptor (TCR) chains and detecting preferential proliferation of a V β subset.---

--57. (Amended) A process [Process] according to claim 56, wherein the cells bearing T-cell receptors are T-cell hybridoma bearing defined human V β domains.--

--58. (Amended) A process [Process] for detecting SAg activity of an expressed human retrovirus associated with human autoimmune disease or of a portion of said retrovirus comprising;

- i) transfecting expressed retroviral DNA or portions thereof into MHC Class II⁺ antigen presenting cells under conditions in which the DNA is expressed,
- ii) contacting the transfectants with cells bearing one or more defined (V)- β T-cell receptor chains,

and

- iii) determining whether the transfectant is capable of inducing preferential proliferation of a V β subset, the capacity to induce preferential proliferation being indicative of SAg activity within the transfected DNA or portion thereof.--

--59. (Amended) A process [Process] for isolating and characterising a human retrovirus, particularly a human endogenous retrovirus (HERV), said retrovirus having SAg activity and being involved in human autoimmune disease, comprising the following steps:

- i) isolating [isolation of] the 5' R-U5 ends of expressed putative retroviral genomes using nucleic acid amplification, the 3' primer being complementary to known "primer binding sites" (pbs);
- ii) isolating [isolation of] the 3' R-poly(A) ends corresponding to the 5' R-U5 ends, by use of primers specific for the R regions isolated in step i);
- iii) amplifying [amplification of] the conserved RT-RNase H region within the pol gene by using degenerate primers corresponding to the conserved region;
- iv) amplifying [amplification of] the 5' moiety of the putative retroviral genome by using primers specific for the different U5 regions isolated in step i) in conjunction with a primer specific for the 3' end of the central pol region isolated in step iii);
- v) amplifying [amplification of] the 3' moiety of the putative retroviral genome using primers specific for the central pol region isolated in

09/27/98
A3
enit